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1. A compound of the general formula:

$$R_{a}$$

$$Z'$$

$$R_{0}$$

$$Z''$$

$$R_{0}$$

$$R_{h1}$$

$$R_{h2}$$

wherein:

- a) R_b and R_o are independently -H, -Cl, -Br, -I, -F, -CN, lower alkyl, -OH, -CH2-OH, -NH2; or N(R6)(R7), wherein R6 and R7 are independently hydrogen or an alkyl or branched alkyl with up to 6 carbons;
- b) R_a is -N3, -C \equiv N, -C \equiv C R, -C=CH-R, -R-C=CH2, -C=CH3 -O-R, -R-R1, or -O-R-R1 where R is a straight or branched alkyl with up to 10 carbons or aralkyl, and R1 is -OH, -NH2, -Cl, -Br, -I, -F or CF3;
- c) Z' is >CH, >COH, or >C-R2-OH, where R2 is an alkyl or branched alkyl with up to 10 carbons or aralkyl;
- d) >C-Rg is >CH2, >C(H)-OH, >C=O, >C=N-OH, >C(R3)OH, >C=N-OR3, >C(H)-NH2, >C(H)-NHR3, >C(H)-NR3R4, or >C(H)-C(O)-R3, where each R3 and R4 is independently an alkyl or branched alkyl with up to 10 carbons or aralkyl;
- e) R_{h1} and R_{h2} are independently H, or a straight or branched chain alkyl, alkenyl or alkynyl with up to 6 carbons that is

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unsubstituted, or substituted with one or more groups selected from a hetero functionality (O-Y, N-Y or S-Y) where Y is H, Me or an alkyl chain up to 6 carbons; a halo functionality (F, Cl, Br or 1); an aromatic group optionally substituted with hetero, halo or alkyl; or $R_{\rm h1}$ and $R_{\rm h2}$ are independently an aromatic group optionally substituted with hetero, halo or alkyl, provided that both $R_{\rm h1}$ and $R_{\rm h2}$ are not H:

f) Z" is >CH₂, >C=O, >C(H)-OH, >C=N-OH, >C=N-OR₅, > C(H)-C \equiv N, or >C(H)-NR₅R₅, wherein each R₅ is independently hydrogen, an alkyl or branched alkyl with up to 10 carbons or aralkyl;

and wherein all monosubstituted substituents have either an α or β configuration.

- The compound of Claim 1, wherein:
 R_b and R₀ are H,
 R_a is OCH3
 Z' is >C-OH,
 >C-R_g is >C(H)-β-OH, and
 Z" is >CH2.
- 3. The compound of Claim 2, wherein : R_{h1} and R_{h2} are independently H and Et.
- The compound of Claim 2, wherein:
 R_{h1} and R_{h2} are independently H and n-Pr.
- The compound of Claim 2, wherein:
 R_{h1} and R_{h2} are independently H and i-Bu.
- 6. The compound of Claim 2, wherein:

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8. The compound of Claim 2, wherein : R_{b1} and R_{b2} are independently H and Me.

The compound of Claim 2, wherein:

R_{h1} and R_{h2} are independently H and n-Bu.

- The compound of Claim 2, wherein:
 R_{h1} and R_{h2} are independently H and (CH₂)_n-C(Me)₂.
- 10. A method of inhibiting angiogenesis comprising administering to an endothelial cell an angiogenesis inhibiting amount of a compound of the general formula:

- wherein:
- a) R_b and R_0 are independently -H, -Cl, -Br, -I, -F, -CN, lower alkyl, -OH, -CH2-OH, -NH2; or N(R6)(R7), wherein R6 and R7 are independently hydrogen or an alkyl or branched alkyl with up to 6 carbons:
- b) R_a is -N₃, -C = N, -C = C-R, -C=CH-R, -R-C=CH₂, -C=CH₃. -O-R, -R-R₁, or -O-R-R₁ where R is a straight or branched alkyl with up to 10 carbons or aralkyl, and R₁ is -OH, -NH₂, -Cl, -Br, -I, -F or CF₃;

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- c) Z' is >CH, >COH, or >C-R2-OH, where R2 is an alkyl or branched alkyl with up to 10 carbons or aralkyl;
- d) >C-Rg is >CH2, >C(H)-OH, >C=O, >C=N-OH, >C(R3)OH, >C=N-OR3, >C(H)-NH2, >C(H)-NHR3, >C(H)-NR3R4, or >C(H)-C(O)-R3, where each R3 and R4 is independently an alkyl or branched alkyl with up to 10 carbons or aralkyl;
- e) R_{h1} and R_{h2} are independently H, or a straight or branched chain alkyl, alkenyl or alkynyl with up to 6 carbons that is unsubstituted, or substituted with one or more groups selected from a hetero functionality (O-Y, N-Y or S-Y) where Y is H, Me or an alkyl chain up to 6 carbons; a halo functionality (F, Cl, Br or I); an aromatic group optionally substituted with hetero, halo or alkyl; or R_{h1} and R_{h2} are independently an aromatic group optionally substituted with hetero, halo or alkyl, provided that both R_{h1} and R_{h2} are not H;
- f) Z" is >CH₂, >C=O, >C(H)-OH, >C=N-OH, >C=N-OR₅, >C(H)-C \equiv N, or >C(H)-NR₅R₅, wherein each R₅ is independently hydrogen, an alkyl or branched alkyl with up to 10 carbons or aralkyl;

and wherein all monosubstituted substituents have either an α or β configuration.